

Cambridge International Examinations

Cambridge International Advanced Subsidiary and Advanced Level

CANDIDATE NAME				
CENTRE NUMBER		CANDIDAT NUMBER	E	

BIOLOGY 9700/22

Paper 2 Structured Questions AS

October/November 2015
1 hour 15 minutes

Candidates answer on the Question Paper.

No Additional Materials are required.

READ THESE INSTRUCTIONS FIRST

Write your Centre number, candidate number and name in the spaces provided at the top of this page. Write in dark blue or black pen.

You may use an HB pencil for any diagrams or graphs.

Do not use staples, paper clips, glue or correction fluid.

DO NOT WRITE IN ANY BARCODES.

Answer all questions.

Electronic calculators may be used.

You may lose marks if you do not show your working or if you do not use appropriate units.

At the end of the examination, fasten all your work securely together.

The number of marks is given in brackets [] at the end of each question or part question.

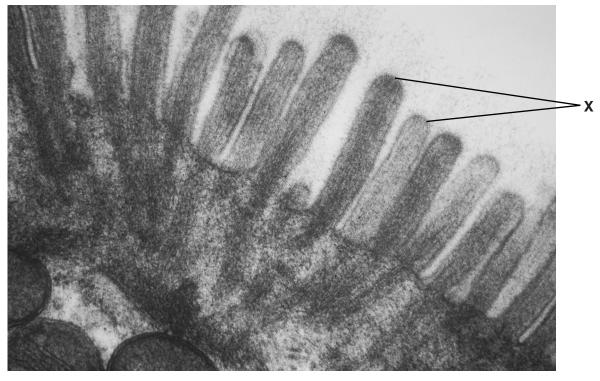
This document consists of 16 printed pages.



Answer all the questions

		nt compared an in e magnification.	nage of a plant co	ell with an imag	e of an animal o	cell. Both image	es were at
Par	ts (a)	to (c) are four co	orrect comparativ	e statements al	oout these imag	es.	
(a)	mei	h cells contain la mbrane of these o always appear to	organelles is fold	ed. These orga			
	(i)	State the name	of the organelles	described.			
							[1]
	(ii)	Suggest one reashape.	ason why the orga	anelles describe	ed do not alway	s seem to have	the same
							[1]
(b)		h cells contain cel a membrane.	ll structures that a	are approximate	ely 25 nm in dia	meter and are r	not bound
	(i)	State the name	of the cell structu	res described.			
							[1]
	(ii)	Draw a circle ard	ound the measure	ement that has	the same value	as 25 nm.	
		0.00025 μm	0.0025 μm	0.025 μm	0. 25 μm	2.5 μm	
							[1]
(c)		ere are strands of ese are not visible			annels in the c	ell wall of the	plant cell.
	(i)	State the name	of the cell structu	res described.			
							[1]
	(ii)	Explain one adv	antage to the pla	ınt cell of havinç	these structur	es.	

(d) Fig. 1.1 is a transmission electron micrograph of part of an epithelial cell from the small intestine of a mammal.



magnification × 65000

Fig. 1.1

me the cell structures labelled X in Fig. 1.1 and state their function.	
	[1]
	[Total: 7]

2

Tob	acco smoking is known to be associated with atherosclerosis and emphysema.
(a)	Outline ways in which tobacco smoking can contribute to atherosclerosis.
	[3
	damaged area of right lung
	Fig. 2.1

State the general role shared by macrophages and neutrophils.
[4]

	(ii)	Suggest how the loss of the elastic fibres would cause the enlargement of the lung shown in Fig. 2.1.
		[2]
(c)	impo	synthesis and release of elastase enzymes by macrophages and neutrophils is an ortant feature in the development and progression of emphysema. Elastase causes the akdown of the protein elastin, the main component of elastic fibres.
	(i)	Explain what is meant by an enzyme.
		[2]
	(ii)	Elastase has an active site with a specific shape. The mode of action of this enzyme supports the lock and key hypothesis.
		Explain the mode of action of elastase.
		You may use the space below to draw a diagram or diagrams to help your answer.
		হে)

- (d) There are two inhibitors of elastase that are produced in the body, TIMP-1 and A1AT:
 - macrophage elastase is inhibited by TIMP-1
 - neutrophil elastase is inhibited by A1AT.

The inhibitors can be inactivated by the elastase enzymes:

- macrophage elastase can inactivate A1AT
- neutrophil elastase can inactivate TIMP-1.

In healthy lungs, the activity of elastase enzymes is regulated. Tobacco smoke can disrupt this regulation.

(1)	inhibitor.
	Suggest how structural changes to A1AT will affect its mode of action.
	[1]
(ii)	A1AT is a protein. Some non-smokers have a mutation in the gene coding for A1AT and are at risk of developing emphysema as there is a lack of A1AT in the lung tissue.
	Explain why a lack of A1AT in these non-smokers means that they are at risk of developing emphysema.
	[3]

(e)	Tobacco smoke is known to cause increased production of macrophage elastase. <i>MMP12</i> is the gene coding for macrophage elastase. Copies of this gene are produced as messenger RNA (mRNA).
	Describe how this mRNA is used in translation to produce macrophage elastase.
	[5]
	[Total: 20]

3 The photomicrographs in Fig. 3.1 show stages of the mitotic cell cycle occurring in the root tip of the onion, *Allium sp.* They are all of the same magnification. Stages **A** to **C** are in the correct sequence and stages **K** to **N** are **not** in the correct sequence.

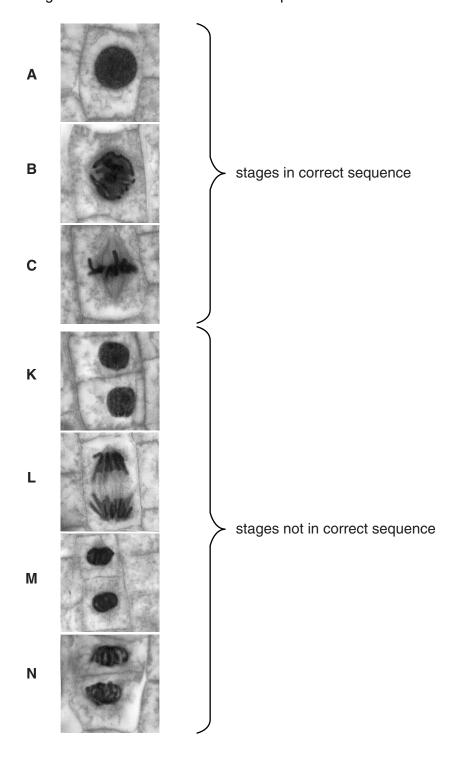


Fig. 3.1

(a)) Name stages A and C.			
	A			
	C			[1]
(b)) Put stages K to N in the correct sequence, stars stage C .	ing with the sta	ge that immediately fo	llows
	$c \longrightarrow $	→	→ 	[1]
(c)) Explain how the behaviour of the chromosomes a that the two daughter cells will be genetically ider	•	g stage L in Fig. 3.1 en	sures
				•••••
			[To	tal: 5]

4 (a) Table 4.1 describes three examples of substances moving into or out of cells.

Complete Table 4.1 by identifying the transport mechanism involved for each example.

Table 4.1

example	transport mechanism involved
uptake of magnesium ions from a lower concentration in the soil solution to a higher concentration in the cytoplasm of a root hair cell	
release of antibodies from an active B-lymphocyte (plasma cell)	
movement of sucrose from a companion cell into a phloem sieve tube element via plasmodesmata	

[3]

(b) Oxygen moves into and out of red blood cells. Fig. 4.1 shows an oxygen dissociation curve for adult human haemoglobin.

percentage saturation of haemoglobin

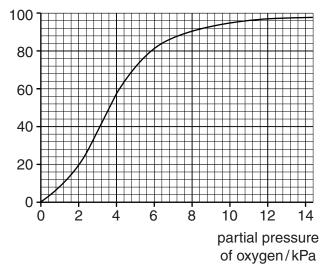


Fig. 4.1

The steepest part of the curve is between 2.6 kPa and 4.2 kPa.

Explain the importance of this for respiring tissues.

.....

[Total: 5]

		11
5 [Dise	eases can be infectious or non-infectious.
((a)	Explain the difference between an infectious and a non-infectious disease.
		[2
		[2
		Malaria is an infectious disease caused by <i>Plasmodium</i> . <i>Plasmodium</i> requires two hosts to complete its complex life cycle. One of the hosts is the <i>Anopheles</i> mosquito, which acts as a vector of malaria.
		Transmission of malaria occurs when females of some species of <i>Anopheles</i> take blood meals from humans infected with <i>Plasmodium</i> , and then feed on uninfected individuals.
		Both male and female <i>Anopheles</i> mosquitos have piercing and sucking mouthparts. The female mosquito is shown in Fig. 5.1.
		Fig. 5.1
((b)	The blood meals are a good source of protein for <i>Anopheles</i> for the production of eggs.
		Explain why blood is a good source of protein

[Turn over

(c) Fig. 5.2 shows the global distribution of those species of *Anopheles* that are able to act as hosts for Plasmodium.

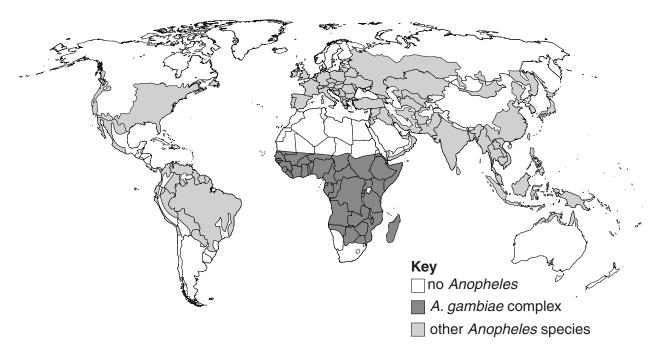


Fig. 5.2

in Fig. 5.2 and the global distribution of malaria.
[3]
[0]

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(i)

(ii)	The distribution of <i>Anopheles</i> shown in Fig. 5.2 includes over forty different species that are vectors of malaria. The areas with the highest number of cases of malaria are also the areas where <i>Anopheles gambiae</i> occurs. <i>A. gambiae</i> is responsible for most of the transmission of the disease in these areas.
	Suggest why A. gambiae is responsible for most of the transmission of Plasmodium.

Fig. 5.3 is part of a complex food web in an area of Kenya where the larvae and adults of *A. gambiae* occur.

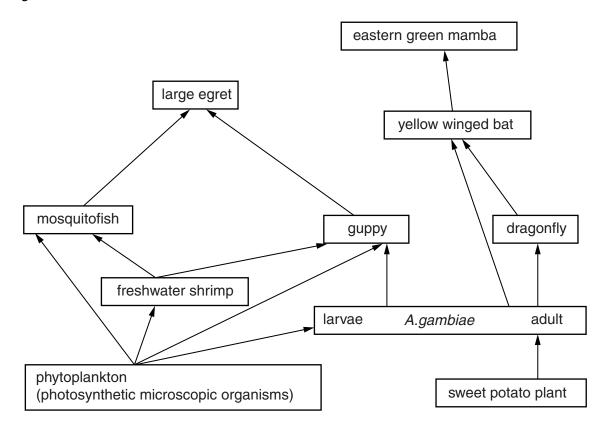


Fig. 5.3

(d)	(i)	Name one organism in Fig. 5.3 that is a tertiary consumer.	
		[1	
((ii)	Explain, in terms of energy transfer, why it is likely that the eastern green mamba feed on other organisms in addition to yellow winged bats.	
		ro	

	(iii)	Suggest how the information in Fig. 5.3 can be used in the control of malaria in other areas of Kenya.
		[2]
(e)		n male and female adult <i>A. gambiae</i> feed on sweet potato plants. Fig. 5.4 shows a sweet to plant.

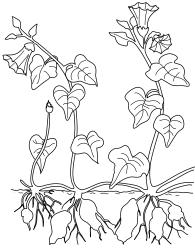


Fig. 5.4

gambiae and e xplain your answer.
[2]

[Total: 18]

6

(a)	The thickness of the different chambers of the mammalian heart is due to the amount cardiac muscle present. The atria have less cardiac muscle than the ventricles, and he thinner walls.						
	In te	erms of their functions, explain why the atria have thinner walls than the ventricles.					
	•••••						
		[2]					
(b)	Nar	ne the dividing wall separating the right and left sides of the mammalian heart.					
		[1]					
(c)	Transpiration and translocation are both processes occurring in plants.						
	(i)	State one way in which transpiration differs from translocation.					
		[1]					
	(ii)	State one way in which transpiration and translocation are similar.					
		[1]					
		[Total: 5]					

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